


Gelatin Tannate for Acute Childhood Gastroenteritis: A Randomized, Single-Blind Controlled Trial

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Abstract

Background Oral rehydration therapy is the recommended treatment for acute childhood gastroenteritis. The aim of this study was to assess the efficacy and safety of gelatin tannate plus oral rehydration compared with oral rehydration alone.

Methods We conducted a multicenter, parallel, randomized, controlled, single-blind, prospective, open-label trial. A central randomization center used computer generated tables to allocate treatments. The study was performed in two medical centers in Italy. Sixty patients 3–72 months of

age with acute gastroenteritis were recruited (median age 18 months; age range 3–66 months): 29 received an oral rehydration solution (ORS) and 31 an ORS plus gelatin tannate (ORS + G). The primary outcome was the number of bowel movements 48 and 72 h after initiating treatment. Secondary outcomes were: duration of diarrhea, stool characteristics and adverse events.

Results No patient was lost at follow-up. No significant difference in the number of bowel movements after 48 h was reported (2.7 ± 1.3 ORS + G; 3.2 ± 0.8 ORS; $p = 0.06$), although the ORS + G group showed a significant improvement in stool consistency (3.7 ± 1.0 vs. 4.3 ± 0.8 ; $p = 0.005$). At 72 h, a significant reduction in bowel movements was reported in the ORS + G group compared with the ORS group (1.0 ± 1.4 vs. 2.0 ± 1.7 ; $p = 0.01$). Mean duration of diarrhea was significantly lower in the ORS + G group than in the ORS only group (76.8 ± 19.2 vs. 108 ± 24.0 h; $p < 0.0001$). No adverse events were reported.

Conclusions Gelatin tannate added to oral rehydration in children with acute diarrhea was associated with a significant decrease in bowel movements at 72 h, with an early improvement in the stool consistency and shorter disease duration.

Clinical Trial Registration NCT02644200—Gelatin Tannate as Treatment for Acute Childhood Gastroenteritis (<https://www.clinicaltrials.gov>).

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Key Points

Oral rehydration therapy is the main treatment recommended for acute gastroenteritis. Recently, gelatin tannate has been proposed as a theoretically effective new treatment. Clinical data on its efficacy are very limited.

This is the first single-blind, prospective, randomized, parallel study in children that demonstrates that gelatin tannate can be effective, well-tolerated, and safe in addition to oral rehydration therapy.

1 Background

Acute gastroenteritis is one of the most common diseases in childhood, with up to 3–5 billion cases of acute gastroenteritis and nearly 2 million deaths per year in children under 5 years. It is also a significant healthcare burden both in developing and developed countries [1]. According to the European Society for Paediatric Gastroenterology, Hepatology and Nutrition (ESPGHAN) guidelines for the treatment of acute gastroenteritis, oral rehydration with hypo-osmolar solutions is the mainstay of therapy [2]. However, although rehydration therapy has achieved a significant reduction in mortality and morbidity due to dehydration, it does not affect the duration of diarrhea [3, 4]. Therefore, active treatments that may impact on the severity of diarrhea have been implemented, including specific probiotics, such as *Lactobacillus GG* or *Saccharomyces boulardii* or drugs such as racecadotril or diosmectite [5–7].

Recently, gelatin tannate, a compound based on tannic acid suspension in a gelatin solution, has been proposed as a theoretically effective treatment for pediatric acute gastroenteritis [8]. Tannins are mostly water-soluble phenolic compounds, and are well-known antioxidants sourced from medicinal plants, foods, and edible fruits. Vegetable tannins have attracted a lot of attention in recent years because of their multifunctional properties, beneficial to human health, demonstrated through in vitro and in vivo assays, and recently, an antidiarrheic effect of gelatin tannate has been reported [9, 10]. Scaldaferrì et al. evaluated the therapeutic effect and mechanisms of action of gelatin tannate using the dextran sulfate sodium (DSS)-induced acute colitis mouse model. Gelatin tannate not only significantly improved disease severity in terms of disease activity index (DAI), but also body weight and histological score [11].

Furthermore, gelatin tannate has been reported to modulate the composition of gut microbiota, restore mucus layer, with an improvement of gut permeability, and exert a direct anti-inflammatory effect in the gastrointestinal mucosa [12, 13]. Clinical data on the efficacy of gelatin tannate in acute gastroenteritis are very limited. In 2009, Esteban Carretero et al. reported a significant reduction in the number of loose stools in 97 children treated with oral rehydration solution (ORS) plus gelatin tannate (ORS + G), compared with 114 treated with ORS alone in a non-randomized, non-blinded clinical study [10]. No drug-related adverse events were reported. More recently, a randomized, blinded, placebo-controlled trial was performed in 40 adults with acute diarrhea [14]. A significant reduction in the number of loose stools was reported in the active group compared with placebo. Based on these two studies, a systematic review on the efficacy of gelatin tannate in acute gastroenteritis published in 2014 concluded that there is not enough evidence to support the use of gelatin tannate in clinical practice [15].

The aim of this study was to evaluate the efficacy and safety of gelatin tannate combined with ORS compared with ORS in children with acute gastroenteritis.

2 Methods

This was a single-blind, prospective, randomized, parallel study conducted in two pediatric services of two tertiary-care hospitals.

Children were eligible for the study if they were 3–72 months of age with acute gastroenteritis, as defined by having at least three loose stools within the previous 24 h and/or a change in stool consistency to loose or liquid according to the modified Bristol Stool Form Scale for Children (m-BSFS-C) lasting for no longer than 3 days [16, 17]. Patients with gastroenteritis for more than 3 days, those with chronic gastrointestinal diseases, or those receiving other antidiarrheal drugs within 2 weeks prior to enrollment (i.e., antibiotics, probiotics, salicylates, loperamide, racecadotril, diosmectite) were excluded from the study.

All children were randomly assigned to the groups using a computer-generated randomization list. The sequence was concealed and envelopes for consecutive patients prepared accordingly. As each patient was recruited, the next envelope in sequence was opened to determine the allocation of the patient. Group 1 received ORS (controls) and group 2 received ORS plus gelatin tannate (Gelen-terum sachet 250 mg, manufactured by Novintethical Pharma Sagl, Lugano, Switzerland, marketed by ACRAF S.p.A Italy) as follows: 250 mg every 6 h if weighing less than 15 kg and 500 mg every 6 h if weighing over 15 kg,

for 5 days. Allocation and management of enrolled patients were performed by two physicians (M.M. and C.T.), while the study coordinator (M.A.) was blinded to the group allocation and the treatment received until the completion of the trial.

Patients in both treatment groups received a standard rehydration solution, following the recommendations of the ESPGHAN [2].

During the initial visit, the following data were collected: demographic variables, family history of gastrointestinal disease, patient's medical history and characteristics of the current gastroenteritis. Physical examination was performed, collecting data on weight, body temperature, heart and respiration rates, blood pressure, degree of dehydration, and any additional tests required for clinical purposes by the treating physician. Dehydration was evaluated using the Clinical Dehydration Scale (CDS), assessing four items: general appearance, eyes, mucous membranes, and tears [18]. Parents received a symptom diary (see electronic supplementary material, online resource 1) and were instructed to complete it, assessing the number of stools, stool consistency according to m-BSFS-C, food and liquid intake, adverse events, and need for other therapies during the study period. All parents were also instructed to determine the daily number of stools. During the 24-, 48-, and 72-h evaluation, tolerance to treatment, side effects, changes in concomitant medication, need for additional visits to the pediatrician or emergency room, as well as data regarding the outcome of the gastroenteritis (number of bowel movements per day and consistency, vomiting, and weight) were monitored. The duration of diarrhea was defined as the time in hours from enrollment to the last abnormal (loose or liquid) stool. Last abnormal stool was defined when the child passed normal stool or no stool for the next 24 h. Stool consistency was evaluated on m-BSFS-C and defined as follows: (1) separate hard lumps, like nuts; (2) sausage-shaped but lumpy; (3) like a sausage or snake, smooth and soft; (4) fluffy pieces with ragged edges, a mushy stool; and (5) watery, no solid pieces [16]. Therapeutic adherence was evaluated on the basis of the number of doses received in 48 h of treatment; good compliance was defined as when the patients received 75% or more of doses according to the patient's bodyweight.

At the last visit, on day 7, symptom diary, including side effect occurrence, was evaluated.

The main objective of this study was to determine the efficacy of treatment according to the number of bowel movements after 48 and 72 h after initiation of therapy with ORS + G versus ORS alone. Secondary aims were to evaluate differences in the length of the diarrhea, stool characteristics, and the safety and tolerance to the study drug.

The study protocol was defined in accordance with the Declaration of Helsinki, and was approved by the ethical committee of each participating center (593/12). Written informed consent was obtained from parents of all children.

2.1 Statistical Analysis

All data were summarized and displayed as mean \pm standard deviation (SD) for the continuous variables. Categorical data were expressed as frequencies and percentages. Comparison of groups was performed using Student's *t* test for unpaired data in a two-group comparison and one-way analysis of variance (ANOVA) with Bonferroni's test for multiple group comparisons. Chi-square tests with Fisher's correction were used to address any differences for categorical variables, as needed. A *p* value of 0.05 or less was considered as significant. Assessment of the duration of diarrhea and specific variables was performed using Pearson and Spearman rank correlation. Kaplan–Meier survival analysis was used to estimate the survival distribution of the duration of diarrhea in the two groups. GraphPad statistical package was used to perform all statistical analyses (GraphPad Software, Inc.).

3 Results

Ninety-six children (42 males; median age 19.6 months, age range 3–69 months) were initially recruited from June 2012 to June 2014: three were excluded because of concomitant chronic diseases; 19 were already receiving racecadotril; one started antibiotic therapy; two who received ORS needed intravenous rehydration; and 11 refused to take part in the study. Therefore, 60 children [30 males (50%); median age 18 months, age range 3–66 months] were included in the study (Fig. 1).

Patients were randomized as follows: 29 (median age 18 months; range 7–43 months) received an ORS; 31 children (median age 20 months; range 3–66 months) received ORS + G.

Both groups were similar in terms of baseline clinical characteristics: median age, gender distribution, weight, concomitant symptoms, and CDS (Table 1). Seventeen patients in group 1 and 14 in group 2 self-managed their toileting.

The number of bowel movements at enrollment was 5.1 ± 2.5 in the ORS group and 6.2 ± 2.9 in the ORS + G group ($p = 0.11$).

At 24 and 48 h, the number of bowel movements was not significantly different between the two groups (3.0 ± 1.1 ORS vs. 2.9 ± 1.0 ORS + G; $p = 0.81$ and 3.2 ± 0.8 vs. 2.7 ± 1.3 ; $p = 0.06$, respectively). However, at 72 h, children with ORS + G had a significant reduction

Fig. 1 Consort flow diagram of the two groups' progress through the phases of the parallel randomized trial: enrollment, intervention allocation, follow-up, and data analysis

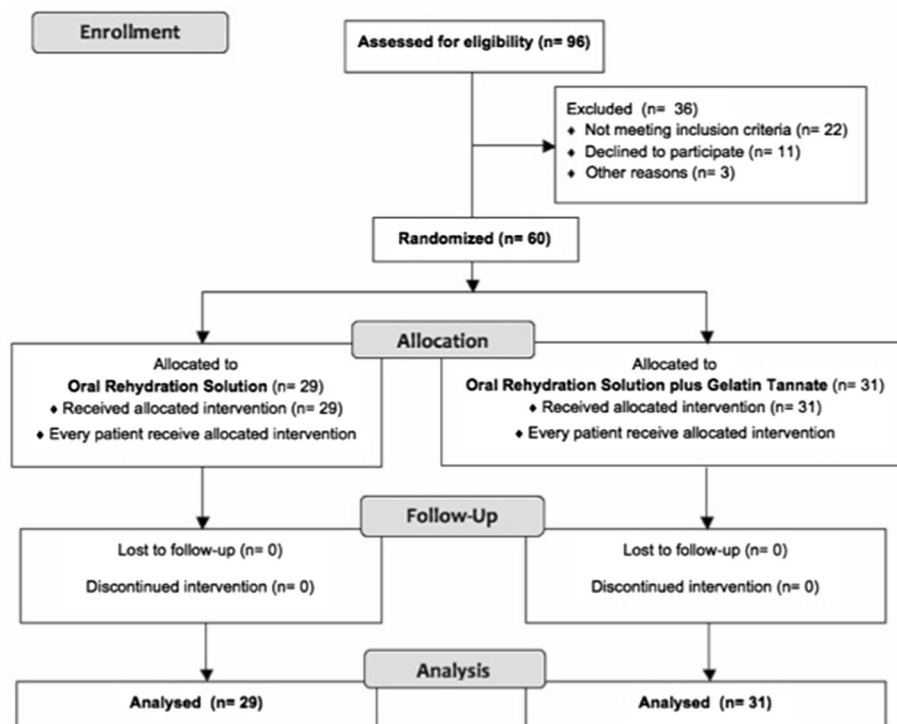


Table 1 Baseline clinical characteristics of the children enrolled in the study

	ORS group (<i>N</i> = 29)	ORS + G group (<i>N</i> = 31)	<i>p</i>
Median age (range), months	18 (7–43)	20 (3–66)	0.14
Male (%)	15 (52)	15 (48)	1.0
Mean weight (kg ± SD)	10.77 ± 2.31	12.02 ± 3.27	0.09
Mean blood pressure (mmHg)	Systolic: 102.3 ± 6.97 Diastolic: 58.9 ± 3.14	Systolic: 104.5 ± 9.02 Diastolic: 62.5 ± 5.22	0.86 0.77
Duration of diarrhea before enrollment (h)	79.9	55.0	0.12
Bowel movements/day	5.13	6.29	0.11
Abdominal pain (%)	8 (28)	15 (48)	0.11
Vomiting (%)	21 (72)	16 (52)	0.11
Fever (%)	18 (62)	14 (45)	0.20
Mean Clinical Dehydration Scale	3.10	3.26	
Dehydration (%)	Mild: 3 (10) Moderate: 25 (86) Severe: 1 (4)	Mild: 2 (6) Moderate: 26 (84) Severe: 3 (10)	1.0 0.47 0.35

ORS oral rehydration solution, ORS + G oral rehydration solution plus gelatin tannate, SD standard deviation

in bowel movements (1.0 ± 1.4) compared with those treated with ORS (2.0 ± 1.8 ; $p = 0.01$) (Fig. 2). At 48 h, a significant improvement in bowel consistency, according to m-BSFS-C, was reported in the ORS + G group compared with the ORS group (3.7 ± 1.0 and 4.3 ± 0.8 ; $p = 0.005$).

The mean duration of diarrhea was significantly shorter in patients treated with ORS + G than in those treated with ORS only (76.8 ± 19.2 vs. 108 ± 24.0 h; $p < 0.0001$), with a mean difference of 31.2 h. Figure 3 shows the time

to diarrhea resolution in the two groups (log-rank 5.7; $p = 0.01$). In a univariate analysis of baseline clinical variables related to the duration of diarrhea at 72 h, we found age to be the only factor correlated [$r = 0.26$; 95% confidence interval (CI) 0.01–0.48; $p = 0.03$] (Table 2). After 72 h of treatment, 28 patients (96.5%) in the ORS group and 29 (93.5%) in the ORS + G group did not complain of any symptoms: one child (3%) in the ORS group and one (3%) in the ORS + G complained of

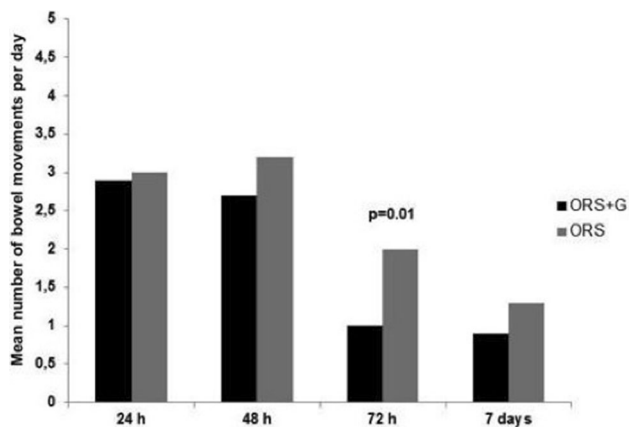


Fig. 2 Mean of bowel movements per day. ORS + G children had a significant reduction in the bowel movements compared with those treated with ORS only at 72 h. ORS oral rehydration solution, ORS + G oral rehydration solution plus gelatin tannate

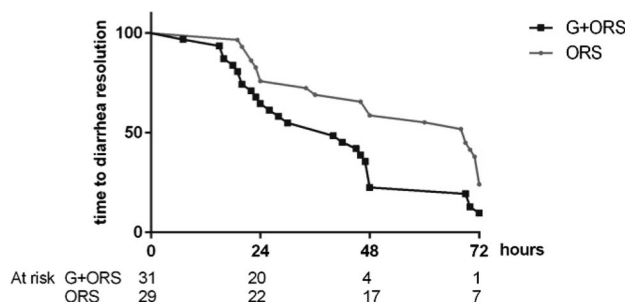


Fig. 3 Kaplan–Meier analysis. The Kaplan–Meier estimator shows the fraction of patients still affected by diarrhea after the start of treatment. ORS oral rehydration solution, G + ORS oral rehydration solution plus gelatin tannate

abdominal pain, and one in the ORS + G group of vomiting. However, diarrhea was still present in seven patients of the ORS group (24.1%) and in one patient of the ORS + G group (3.2%).

Three patients in the ORS group and one in the ORS + G group needed additional visits to the primary doctor during treatment and one of the ORS group resorted to the emergency room; all had diarrhea lasting more than 72 h.

Table 2 Univariate analysis of baseline clinical factors related to the persistence of diarrhea after 3 days in 60 children with acute gastroenteritis

	<i>r</i>	95% CI	<i>P</i> values
Age	0.26	0.01 to 0.48	0.03
Gender (male)	0.06	−0.19 to 0.31	0.60
Symptom duration at the enrollment	0.06	−0.19 to 0.30	0.64
Number of bowel movements	−0.02	−0.27 to 0.23	0.85
Vomiting	−0.15	−0.39 to 0.1	0.23
Abdominal pain	0.20	−0.05 to 0.43	0.11
Fever	0.11	−0.14 to 0.36	0.37

CI confidence interval

All parents filled out the diary completely.

At 7-day follow-up, tolerability and drug acceptance were evaluated: all patients received 75% or more of the drug doses according to the patient's weight. One patient reported the occurrence of nausea about 20 min after administration of gelatin tannate. No other adverse events were reported.

4 Discussion

The most recent ESPGHAN guidelines for the treatment of acute gastroenteritis clearly state that oral rehydration with a hypo-osmolar solution is the main treatment and should be started as soon as possible [2].

Recent data have highlighted and introduced the possibility of using gelatin tannate, a mucosal barrier protector, as an innovative approach in the management of intestinal diseases, with the aim of enhancing mucus barrier activity and restoring the gut barrier [11]. Nevertheless, there are very limited data on its efficacy in children with acute diarrhea, although a randomized controlled trial performed in adults with acute gastroenteritis reported its good efficacy and safety in this setting [14].

The objective of our single-blind, prospective, randomized, parallel study was to compare the efficacy of combined therapy with gelatin tannate and oral rehydration with that of oral rehydration alone in children younger than 6 years affected with acute gastroenteritis as outpatient care.

We found patients treated with ORS + G to have a significant decrease in the number of bowel movements after 72 h of treatment, when compared with those managed with ORS alone, while no significant difference was shown after 24 and 48 h, and a significant improvement of bowel consistency was seen after 48 h of treatment.

Since it has already been demonstrated that gelatin tannate exerts its action by restoring the physiological function of the intestine, one may speculate that the drug is able to provoke a clinical effect only after restoring the intestinal barrier and thus will not be immediately effective [9, 11]. Furthermore, the slow action may be also due to the well-

known astringent properties of tannins allowing the absorption of proinflammatory mucoproteins from the intestinal mucus and their elimination through the feces [19].

Overall, the mean duration of diarrhea was significantly shorter in patients treated with ORS + G than in those given ORS only, with a mean difference of 31.2 h. These data are in keeping with the clinical effect of gelatin tannate in adults and with previous studies on the efficacy of other drugs suggested for the treatment of acute gastroenteritis [14]. Salazar-Lindo et al. reported a reduction in the duration of diarrhea in racecadotril-treated children up to 44 h, compared with controls, while diosmectite is effective in reducing acute gastroenteritis by 23.39 h [7, 20].

Compliance with administration was good; every patient received 75% or more of doses according to the patient's weight. This result confirms a good ease of administration. Safety was similar for the two treatment groups, with only one patient reporting the occurrence of nausea during a single administration of gelatin tannate. Three patients in the ORS group and one in the ORS + G group needed additional visits to the primary doctor during treatment, and one in the ORS group needed additional evaluation at the emergency room. These data suggest that duration of illness could be an important reason to utilize primary care or the emergency room.

We tried to identify predictive factors for prolonged diarrhea, and we found only age to be significantly correlated, with a duration of diarrhea greater than 3 days for older compared with younger children. Conversely, a previous study conducted in developing countries reported young age to be a risk factor for prolonged diarrhea [21]. This result could be related to the different study setting and design and should be confirmed in larger trials.

The strengths of our study include adequate randomization and the use of intention-to-treat analysis, both of which minimize the risk of bias. However, there are several limitations. First of all, the small sample size. Baseline characteristics were similar in the two groups of patients, and patients were randomized according to a computer-generated protocol with a single-blind design in order to minimize the risk of bias.

Second, clinical measures (i.e., number of bowel movements and duration of diarrhea) were used as outcomes. These measures alone are not considered optimal, indeed stool output is the main criterion defined by the World Health Organization to conclude that a drug is effective in acute diarrhea [22]. Since gelatin tannate was used as outpatient therapy, we could not assess the 24 h stool output, and we decided to evaluate clinical outcomes of efficacy. We evaluated patients at 24, 48 and 72 h, and we made an additional visit at day 7, in order to obtain accurate data, and we used a symptom diary for the entire duration of the study. Furthermore, we monitored common

clinical symptoms such as fever, vomiting, abdominal pain, development of any other new symptoms and any hypersensitivity reaction like skin rashes to define potential adverse effects of intervention.

In contrast to other studies, we did not perform stool cultures at baseline [20, 23]. Nevertheless, ESPGHAN does not recommend performing stool cultures for acute gastroenteritis in primary healthcare, and we did not aim to evaluate the impact of gelatin tannate based on the different etiologies of diarrhea [2]. Finally, in our study design, patients were not hospitalized, the control group did not receive a placebo, and there was no direct monitoring of drug administration.

5 Conclusions

In conclusion, the results of this study suggest that gelatin tannate could be effective, well-tolerated, and safe in addition to oral rehydration therapy in infants and children with acute diarrhea. Further randomized, double-blind, placebo-controlled studies are needed to confirm its efficacy and safety and to evaluate its routine use in children with acute gastroenteritis.

Compliance with Ethical Standards

The study protocol was defined in accordance with the Declaration of Helsinki, and was approved by the ethical committee of each participating center (number of protocol 593/12). Written informed consent was obtained from parents of all children.

Conflict of interest Maurizio Mennini M.D., Carlo Tolone, M.D., Antonella Frassanito M.D., Fabio Midulla M.D. Ph.D., Salvatore Cucchiara M.D. Ph.D. and Marina Aloï M.D., Ph.D. declare that they have no conflict of interest and no financial relationships relevant to this article to disclose.

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Contributors' statement Dr. Aloï and Dr. Mennini conceptualized and designed the study, drafted the initial manuscript, and approved the final manuscript as submitted. Dr. Tolone and Dr. Mennini allocated and managed enrolled patients. Dr. Aloï and Dr. Frassanito designed the data collection instruments and carried out the initial analyses. Prof. Cucchiara and Prof. Midulla critically reviewed the manuscript and approved the final manuscript as submitted.

Availability of data and materials Data will not be shared, because of institutional policy. We will, however, be available to share them in the event of a specific request from a reviewer.

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